

WHAT IS CLAIMED IS:

1. A method of nucleic acid sequencing comprising the steps:

5 (a) amplifying a nucleic acid sample to produce an amplified DNA product;

(b) extending a sequencing primer bound to the DNA product in the presence of terminating nucleotide analogs to  
10 produce a collection of labeled nucleic acid products;

(c) detecting a total amount of label present in the collection to produce a measurement; and

(d) combining a plurality of measurements to determine  
15 DNA sequence information about the sample.

2. A method as described in Claim 1 wherein each measurement of a label corresponds to an amount of terminating nucleotide.

3. A method as described in Claim 1 wherein the DNA sequence information corresponds to a length of the DNA sequence.

4. A method as described in Claim 1 wherein the DNA sequence  
25 information corresponds to a plurality of bases in the DNA sequence.

5. A method as described in Claim 1, wherein after the combining  
30 step, the DNA sequence information is used for human identification.

6. A method as described in Claim 1, wherein after the combining step, the DNA sequence information is used for diagnostic testing.

FOR 500 600 700 800 900 1000 1100 1200 1300 1400 1500 1600 1700 1800 1900 2000 2100 2200 2300 2400 2500 2600 2700 2800 2900 3000 3100 3200 3300 3400 3500 3600 3700 3800 3900 4000 4100 4200 4300 4400 4500 4600 4700 4800 4900 5000 5100 5200 5300 5400 5500 5600 5700 5800 5900 6000 6100 6200 6300 6400 6500 6600 6700 6800 6900 7000 7100 7200 7300 7400 7500 7600 7700 7800 7900 8000 8100 8200 8300 8400 8500 8600 8700 8800 8900 9000 9100 9200 9300 9400 9500 9600 9700 9800 9900 10000

7. A method as described in Claim 1, wherein after the combining step, the DNA sequence information is used for genetic localization or gene discovery.

5 8. A method as described in Claim 1, wherein after the combining step, the DNA sequence information is used for criminal justice applications.

10 9. A method as described in Claim 1, wherein after the combining step, the DNA sequence information is used in conjunction with a DNA database of genetic polymorphisms.

10. A method as described in Claim 1, wherein after the combining step, the DNA sequence information is used for cancer assessment.

11. A system for nucleic acid sequencing comprising:

(a) a means for amplifying a nucleic acid sample to produce an amplified nucleic acid product;

(b) a means for extending a sequencing primer bound to the DNA product in the presence of terminating nucleotide analogs to produce a collection of labeled nucleic acid products, said extending means in connection with the amplified product;

(c) a means for detecting a total amount of label present in the collection to produce a measurement, said detecting means in connection with the collection; and

(d) a means for combining a plurality of measurements to determine DNA sequence information about the sample, said combining means in connection with the measurement.

12. A system as described in Claim 11, wherein the amplifying means includes a PCR thermocycler, the extending means includes a chamber that permits DNA sequencing reactions to occur in the presence of terminating nucleotide analogs, the detecting means  
5 measures fluorescent or other labels that quantify an amount of DNA molecules, and the combining means includes a computing device with memory.

13. A method for obtaining information about a signal comprising  
10 the steps:

(a) inducing a decay function;

(b) imposing the decay function on a signal;

(c) forming a numerical quantity that characterizes the signal's behavior in the presence of the decay function;

(d) combining a plurality of such numerical quantities to obtain information about the signal.

14. A method as described in Claim 13 wherein the signal is a nucleic acid sequence, the decay function is induced by introducing dideoxy terminator analogs into a sequencing reaction,  
25 the numerical quantities correspond to Laplace transform coefficients, and the obtained information helps characterize the sequence.

15. A method as described in Claim 14 wherein the characterization  
30 does not completely describe the nucleic acid sequence.

16. A method as described in Claim 15 wherein the incomplete sequence information describes a genetic polymorphism.